AMENDMENTS TO THE CLAIMS

1.(Withdrawn, Currently amended) Substituted 6-(2-halogenphenyl) triazolopyrimidines A substituted 6-(2-halogenphenyl)-triazolopyrimidine of formula I

in which

Hal is halogen;

L1, L3 independently denote hydrogen, halogen, or C1-C4-alkyl;

L2 is hydrogen, halogen, C1-C4-haloalkyl, or NH2, NHRb, or N(Rb)2,

R^b is C₁-C₈-alkyl, C₃-C₁₀-alkenyl, C₃-C₁₀-alkynyl, C₁-C₆-haloalkyl, C₃-C₆haloalkenyl, C₃-C₆-haloalkynyl, C₁-C₈-alkoxy-C₁-C₈-alkyl, C₁-C₈-alkylthio-C₁-C₈-alkyl, C₃-C₁₀-evcloalkyl, or C(=O)-A, in which

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A is hydrogen, hydroxy, C₁-C₈-alkyl, C₁-C₈-alkoxy, C₁-C₆halogenalkoxy, C₁-C₈-alkylamino or di-(C₁-C₈-alkyl)amino;

wherein at least one from L1, L2, and L3 is not hydrogen;

- X is halogen, cyano, C₁-C₀-alkyl, C₁-C₀-alkoxy, C₁-C₀-haloalkoxy or €₃-C₀alkenyloxy, C₁-C₀-alkenyloxy;
- $R^{1} \ denote \ C_{1}-C_{10}-alkyl, \ C_{2}-C_{10}-alkenyl, \ C_{2}-C_{10}-alkynyl, \ or \ C_{4}-C_{10}-alkadienyl, \ C_{2}-C_{10}-alkenyl, \ or \ C_{4}-C_{10}-alkadienyl, \ or \ c_{4}-c_{10}-alkadienyl, \ c_{7}-c_{10}-alkenyl, \ or \ c_{8}-c_{10}-alkenyl, \ or \ c_{10}-alkenyl, \ or \ c_{10}-a$

wherein R1 may be unsubstituted or may carry one to three groups R4,

- R^a is cyano, nitro, hydroxyl, C₁-C₆-alkyl, C₃-C₆-eyeloalkyl, C₁-C₆-alkoxy, C₁-C₆-alkylthio, C₁-C₆-alkylamino, di-C₁-C₆-alkylamino, C₂-C₆-alkenyl, C₂-C₆-alkynyloxy, C₂-C₆-alkynyloxy; or C₁-C₄-alkylenedioxy;
- R² is hydrogen; and R² is hydrogen.
- (Withdrawn, Currently amended) Compounds The compound of formula I according to claim I, in which

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R¹ is straight chained or branched C₂—C₆—alkenyl, or a straight chained or branched C₁-

Co-alkyl.

3. (Withdrawn, Currently amended) Compounds The compound of formula I according to

claim 1 or 2 in which X is halogen.

4. (Withdrawn, Currently amended) Compounds The compound of formula I according to

claim 1 in which the 6-(2-halogenphenyl)group represents one of the following moleties:

2,3,5-trifluorophenyl, 2 F,4 CF₃ phenyl, 2 F,5 CH₃ phenyl, 2 Cl,4 F phenyl, 2 F,4 Cl

phenyl, 2 F,4 Br phenyl, 2 Cl.4 Br phenyl, 2.3 diffuorophenyl, 2.4 diffuorophenyl, 2.4.5

trifluorophenyl, 2,3,4 trifluorophenyl, 2-F,4-NHC(O)CH₂-phenyl, 2-Br,3,5-

difluorophenyl, 2-F,4-NO2-phenyl, and 2-Cl,4-NO2-phenyl 2.3.5-trifluorophenyl; 2-F,4-

 $\underline{CF_3-phenyl;\ 2-F,5-CH_3-phenyl;\ 2-Cl,4-F-phenyl;\ 2-F,4-Cl-phenyl;\ 2-F,4-Br-phenyl;\ 2-F,4-Br-phenyl;\ 2-F,4-Cl-phenyl;\ 2-F,4-Br-phenyl;\ 2-F,4-Br-p$

Cl.4-Br-phenyl; 2,3-diffuorophenyl; 2,4-diffuorophenyl; 2,4,5-triffuorophenyl; 2,3,4-

trifluorophenyl; 2-F,4-NHC(O)CH₃-phenyl; 2-Br,3,5-difluorophenyl; 2-F,4-NO₂-phenyl;

and 2-Cl,4-NO2-phenyl.

5. (Withdrawn) A process for the preparation of compounds of formula I as defined in claim

3 which comprises reacting 5-amino-1,2,4-triazole

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with 2-phenyl-substituted malonic acid ester of formula II,

wherein Hal, L^1 , L^2 , and L^3 are as defined in formula I, and R denotes C_1 – C_6 -alkyl, under alkaline conditions, to yield compounds of formula III,

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> which are subsequently treated with a halogenating agent to give 5,7-dihalogen-6-phenyltriazoloovrimidines of formula IV

in which Y is halogen, and which is reacted with an amine of formula V

in which \mathbb{R}^1 and \mathbb{R}^2 are as defined in claim 1 to produce compounds of formula I, as defined in claim 1.

6. (Withdrawn, Currently amended) A process for the preparation of compounds of formula I according to claim 1 wherein X is cyano, C₁-C₁₀-alkoxy, or C₁-C₆-haloalkoxy, which comprises reacting 5-halogen-triazolopyrimidine of formula I',

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wherein Y is halogen, with compounds of formula VI,

which are, dependent from the value of X' to be introduced, an anorganic inorganic cyano salt, an alkoxylate, haloalkoxylate or an alkenyloxylate, resp.; respectively, wherein M is ammonium-, tetraalkylammonium-, alkalimetal- or alkaline earth metal cation, to produce compounds of formula I.

(Withdrawn, Currently amended) Intermediates of formulae II, III, and IV An intermediate
of formulae II, III, or IV as defined in claim 5, in which the 6-(2-halogenphenyl)group
represents one of the following moieties:

2,3,5-trifluorophenyl, 2-F,4-CF₂-phenyl, 2-F,5-CH₃-phenyl, 2-Cl,4-F-phenyl, 2-F,4-Cl-phenyl, 2-F,4-Br-phenyl, 2-Cl,4-Br-phenyl, 2-Br,3,5-difluorophenyl, 2-A,5-trifluorophenyl, 2-Br,3,5-difluorophenyl, 2-F,4-NO₂-phenyl, 2-F,4-NO₂-phenyl, 2-Cl,4-Br-phenyl, 2-F,5-CH₃-phenyl, 2-Cl,4-Br-phenyl, 2-F,5-CH₃-phenyl, 2-Cl,4-Br-phenyl, 2-F,4-Cl-phenyl, 2-F,4-Cl-phenyl, 2-F,4-Cl-phenyl, 2-F,4-Cl-phenyl, 2-F,4-Br-phenyl, 2-Cl,4-Br-phenyl, 2-S,4-Br-phenyl, 2-Cl,4-Br-phenyl, 2-B,4-Br-phenyl, 2-Cl,4-Br-phenyl, 2-B,4-Br-phenyl, 2-Cl,4-Br-phenyl, 2-B,4-Br-phenyl, 2-Cl,4-Br-phenyl, 2-B,4-Br-phenyl, 2-

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difluorophenyl; 2-4,5-trifluorophenyl; 2-3,4-trifluorophenyl; 2-F,4-NHC(O)CH;-phenyl; 2-Br,3,5-difluorophenyl; 2-F,4-NO;-phenyl; and 2-Cl,4-NO;-phenyl.

- (Withdrawn) A composition suitable for controlling phytopathogenic fungi, comprising a solid or liquid carrier and a compound of the formula I as claimed in claim 1.
- 9. (Withdrawn) A method for controlling phytopathogenic fungi, which comprises treating the fungi or the materials, plants, the soil or the seed to be protected against fungal attack with an effective amount of a compound of the formula I as claimed in claim 1.
- (Currently amended) Substituted 6-(2-halogenphenyl)-triazolopyrimidines A substituted 6-(2-halogenphenyl)-triazolopyrimidine of formula I

in which

Hal is halogen;

- L1, L3 independently denote hydrogen, halogen, or C1-C4-alkyl;
- L2 is hydrogen, halogen, C₁-C₄-haloalkyl, or NH₂, NHR^b, or N(R^b)₂,
 - R^b is C₁-C₈-alkyl, C₂-C₁₀-alkenyl, C₂-C₁₀-alkynyl, C₄-C₆-haloalkyl, C₃-C₆-haloalkynyl, C₄-C₈ alkyl, C₁-C₈-alkylthio-C₄-C₈-alkyl, C₂-C₆-paloalkyl, or C(≡O)-A, in which
 - A is hydrogen, hydroxy, C₁-C₈-alkyl₇-C₄-C₈-alkoxy, C₄-C₆-halogenalkoxy, C₁-C₈-alkylamino or di (C₁-C₈-alkyl)amino;

wherein at least one from L1, L2, and L3 is not hydrogen;

- X is halogen, eyano, C₁-C₆-alkyl, or C₁-C₆-alkoxy, C₁-C₆-alkoxy, C₁-C₆-haloalkoxy or C₁-C₆-alkenyloxy.
- R¹ and R² together with the interjacent nitrogen atom represent a saturated or partially unsaturated 5- or 6-membered heterocycle, containing one to-four nitrogen atoms or one to three nitrogen atoms and one sulfur or oxygen atom nitrogen atom or one nitrogen atom and one sulfur atom, which ring may be substituted by one to three R⁴ radicals;

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Ra is eyano, nitro, hydroxyl, Ct C6-alkyl, Ct C6-cycloalkyl, Ct C6-alkoxy, Ct-

 $C_6 \text{-alkylthio}, C_4 \cdot C_6 \text{-alkylamino}, \text{di-}C_4 \cdot C_6 \text{-alkylamino}, C_2 \cdot C_6 \text{-alkenyl}, C_2 \cdot C_6 \text{-alkenyl}, C_4 \cdot C_6 \cdot C$

alkenyloxy, C2-C4-alkynyl, C3-C4 alkynyloxy, or C1-C4-alkylonedioxy; is C1-

C6 alkyl.

11. (Currently amended) Compounds The compound of formula I according to claim 10, in

which

R1 and R2 together with the interiacent nitrogen atom represent a heteroevelic ring with 5

or-6 carbon atoms a saturated or partially unsaturated 5- or 6-membered heterocycle.

containing one nitrogen atom or one nitrogen atom and one sulfur atom, being

optionally substituted with one or two C1-C4-alkyl groups.

12. (Currently amended) Compounds The compound of formula I according to claim 10 or 11

in which R1 and R2 together with the interjacent nitrogen atom represent a 5 or 6

membered heterocyclic-ring a saturated or partially unsaturated 5- or 6-membered

heterocycle, containing one nitrogen atom or one nitrogen atom and one sulfur atom, being

optionally substituted with one or two methyl groups.

13. (Currently amended) Compounds The compound of formula I according to claim 10 in

which X is halogen.

 (Currently amended) Compounds The compound of formula I according to claim 10 in which the 6-(2-halogenphenyl)group represents one of the following moieties:

2,3,5 trifluorophenyl, 2 F,4 CF₂-phenyl, 2 F,5 CH₃-phenyl, 2 Cl,4 F-phenyl, 2 F,4 Cl-phenyl, 2 F,4 Br-phenyl, 2 Cl,4 Br-phenyl, 2,3 difluorophenyl, 2,4 difluorophenyl, 2,4,5 trifluorophenyl, 2,3,4 trifluorophenyl, 2 F,4 NIC(O)CH₃-phenyl, 2 Br₂,5 difluorophenyl, 2 F,4 NO₂-phenyl, and 2 Cl,4 NO₂-phenyl 2,3,5-trifluorophenyl; 2-F,4 CF₃-phenyl; 2-F,4 Br-phenyl; 2-Cl,4-F-phenyl; 2-F,4-Cl-phenyl; 2-F,4-Br-phenyl; 2-Cl,4-Br-phenyl; 2,4-difluorophenyl; 2,4,5-trifluorophenyl; 2,3,4-trifluorophenyl; 2,5-trifluorophenyl; 2,3,4-trifluorophenyl; 2-F,4-NIC(O)CH₃-phenyl; and 2-Br₂,3,5-difluorophenyl; 2,3,5-trifluorophenyl; 2,3,4-trifluorophenyl; 2-F,4-NIC(O)CH₃-phenyl; and 2-Br₂,3,5-difluorophenyl; 2.5-trifluorophenyl; 2,5-trifluorophenyl; 2,5-trif

(Withdrawn, Currently amended) A process for the preparation of compounds of formula I
as defined in claims 13 and 14 the compound of formula I as defined in claim 13 which
comprises reacting 5-amino-1,2,4-triazole

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with 2-phenyl-substituted malonic acid ester of formula II,

wherein Hal, L^1 , L^2 , and L^3 are as defined in formula I, and R denotes C_1 – C_6 -alkyl, under alkaline conditions, to yield compounds of formula III,

which are subsequently treated with a halogenating agent to give 5,7-dihalogen-6-phenyltriazolopyrimidines of formula IV Application No. 10/532,719 Docket No.: 4266-0131PUS1 Amendment dated February 20, 2008

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in which Y is halogen, and which is reacted with an amine of formula V

in which R^1 and R^2 are as defined in claim 10 to produce compounds of formula I, as defined in claim 10 defined in claim 13.

16. (Withdrawn, Currently amended) A process for the preparation of compounds the compound of formula I according to claim 10 wherein X is eyane; C₁-C₁₀-alkoxy, or C₁-C₄-haloalkoxy, which comprises reacting 5-halogen-triazolopyrimidine of formula I',

wherein Y is halogen, with compounds of formula VI,

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M-X' VI

which is an alkoxylate, are, dependent from the value of X' to be introduced, an anorganic eyano salt, an alkoxylate, haloalkoxylate or an alkenyloxylate, resp., wherein M is ammonium-, tetraalkylammonium-, alkalimetal- or alkaline earth metal cation, to produce compounds of formula I.

- (Original) A composition suitable for controlling phytopathogenic fungi, comprising a solid or liquid carrier and a compound of the formula I as claimed in claim 10.
- 18. (Withdrawn) A method for controlling phytopathogenic fungi, which comprises treating the fungi or the materials, plants, the soil or the seed to be protected against fungal attack with an effective amount of a compound of the formula I as claimed in claim 10.